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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/763,793	02/27/2001	Hiromasa Miyaji	766.46	3687
5514 7590 10/31/2008 FITZPATRICK CELLA HARPER & SCINTO 30 ROCKEFELLER PLAZA NEW YORK, NY 10112				
EXAMINER				
SHAFFER, SHULAMITH H				
ART UNIT		PAPER NUMBER		
1647				
MAIL DATE		DELIVERY MODE		
10/31/2008		PAPER		

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary

Application No.

09/763,793

Applicant(s)

MIYAJI ET AL.

Examiner

SHULAMITH H. SHAFER

Art Unit

1647

Period for Reply -- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 01 August 2008.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-12 and 46-48 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-12 and 46-48 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☒ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☒ All b) ☐ Some * c) ☐ None of:
1. ☒ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- 1) ☒ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☒ Information Disclosure Statement(s) (PTO/SB/02)
- 4) ☐ Interview Summary (PTO-413)
- 5) ☐ Notice of Informal Patent Application
- 6) ☐ Other: _____
- Paper No(s)/Mail Date 8/1/08, 9/2/08

Detailed Action

Status of Application, Amendments, And/Or Claims:

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 1 August 2008 has been entered.

Claims 1-12 and 46-48 are pending in the instant application and are under consideration. These are the same claims that were submitted on 27 March 2008 and entered as an after final amendment.

Information Disclosure Statement:

The Information Disclosure statements (IDS) submitted on the 1 August 2008 and 2 September 2008 have been considered. The signed copies are attached.

Priority:

Acknowledgment is made of applicants' claim for foreign priority based on an application filed in Japan on 27 of August 1998. A certified copy of the Japan 10/241248 application as required by 35 U.S.C. 119(b). Applicants have provided a certified translation of 10/241248 with submission of 1 August 2008; therefore, Applicants have perfected priority claim to 27 August 1998, the date of filing of Japan 10/241248.

Withdrawn Objections/Rejections

35 U.S.C. § 112, First Paragraph:

The rejection of claims 2, 5, 6, 8-12, 29, 42, 46 and 48 under 35 U.S.C. 112, first paragraph (scope of enablement), is withdrawn in view of applicants amendments to the claims.

The rejection of Claims 2, 5, 6, 8-12, 29, 42, 46 and 47 under 35 U.S.C. 112, first paragraph, as failing to comply with the written description requirement is withdrawn in view of applicants' amendment to the claims.

Maintained Rejections

35 U.S.C. §§ 101 and 112, First Paragraph:

35 U.S.C. § 101 reads as follows:

Whoever invents or discovers any new and useful process, machine, manufacture, or composition of matter, or any new and useful improvement thereof, may obtain a patent therefor, subject to the conditions and requirements of this title.

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

The rejection of Claim(s) 1-12 and 46-48 under 35 U.S.C. 101 because the claimed invention is not supported by either a credible, substantial or specific asserted utility or a well established utility is maintained for reasons of record and reasons set forth below.

Applicants traverse the rejection (After final submission of 27 March 2008 and Remarks of 1 August 2008).

The reasons for the traversal are:

a. the polypeptide of SEQ ID NO:1 is a member of the nucleoside transporter system. Dipyridamole, a compound known to inhibit the uptake of adenosine, enhances bronchospasm in an asthmatic patient (Crimi et al. 1988 Allergy 43:179-83, submitted with after-final amendment). Thus, one would conclude that bronchospasm is suppressed by accelerating uptake of adenosine; therefore, when a DNA encoding SEQ ID NO:1 is expressed in the lung of asthmatic patients, bronchospasm in the patient is treated (submission of 27 March 2008, page 7, last paragraph, bridging page 8, 1st paragraph).

b. dipyridamole, as an inhibitor of equilibrative nucleoside transporters, including the one described in the instant invention, inhibits the decrease of extracellular adenosine concentration by inhibiting uptake of adenosine. The transporters take in and eliminate extracellular adenosine. Thus, when transporters are inhibited, more adenosine is available in the extracellular space to activate the adenosine receptor on the cell membrane and cause bronchoconstriction (Remarks of 1 August 2008, page 2, numbered paragraphs 1-3).

c. References provided with applicants submission provide evidence that concentration of adenosine is elevated in the lung of asthmatic and adenosine-induced bronchoconstriction is mediated by A(1) receptor on the cell membrane.

Response to Arguments

Applicant's arguments have been fully considered but are not found to be persuasive for the following reasons:

Applicants have sought to establish the following fact pattern:

1. Adenosine, present in elevated levels in the lungs of asthmatic patients, binds to the A(1) receptor, and elicits bronchoconstriction (Hua et al. 2007. American Journal of Physiology-Lung, Cellular and Molecular Physiology. 293:L25-32 and Brown et al. 2008. Eur. Resp J. 31:311-9, abstracts of both references submitted on 1 August 2008).
2. Nucleoside transporters, including the polypeptide of the instant invention, transport adenosine into the cell, thereby reducing the amount of adenosine available to

activate the A(1) receptor; thus these polypeptides play a role in reducing bronchospasms.

3. Therefore, the polypeptide of the instant invention and its encoding DNA would be useful in treating bronchospasm in the asthmatic patient.

In summary, Applicants assert, in above arguments (a, b, 2 and 3 above), that the polypeptide of the instant invention and DNA encoding said polypeptide have utility in treating bronchospasm in an asthmatic patient, for which there is no support in the specification of the instant invention.

Applicants are reminded that a specific or substantial asserted utility or a well established utility must be presented at the time of filing. The specification has not asserted a specific and substantial utility nor is there a well established utility for the claimed invention because the specification and/or the art fail to establish a connection between the polypeptide of SEQ ID NO:1 structure, expression or activity or changes in structure, expression or activity and any specific disease state nor has this been established for the encoding DNA (SEQ ID NO:2). Applicants assert, in the specification, that the polypeptide of the instant invention may be used as "a preventive agent or a therapeutic agent for ischemic heart disease, cerebral disorder at the time of stroke, immune response accompanied by organ transplantation, malignant tumor, nephritis, pancreatitis or hypertension....Its applications as an analgesic, an antiplatelet agent, an agent for increasing activity of an antiviral agent or a malignant tumor treating agent and an agent for reducing side effects at the time of chemotherapy can also be expected" (page 63, last paragraph, bridging page 64, 1st paragraph). There is no assertion that the polypeptide of the instant invention (or its encoding DNA) may be used to treat symptoms in the asthmatic patient.; thus there is no support in the specification for this utility.

In response to a: Applicants argue, without supporting evidence, that when a DNA encoding SEQ ID NO:1 is expressed in the lung of asthmatic patients, bronchospasm in the patient is treated by gene therapy. The specification does not

contemplate any gene therapy methods or protocols and does not support this assertion of utility.

In response to b: One of skill in the art would be unable to predict that nucleoside transporters, such as the polypeptide of the instant invention would have a therapeutic role in treatment of asthma and related bronchospasms. The fact that dipyridamole, an inhibitor of equilibrative nucleoside transporters, enhances bronchospasm in an asthmatic patient does not provide evidence that stimulating nucleoside transporters or increasing the level of nucleoside transporter protein in the cell would inhibit bronchospasms. Contrary to applicants' assertion above, the art teaches that equilibrative transporters can move adenosine bidirectionally across plasma membranes by facilitated diffusion. Adenosine formed intracellularly can be released by bidirectional nucleoside transport processes to activate cell surface receptors (Borgland et al. 1998, Europ. J of Pharm. 346:339-344, abstract and page 339, 2nd column, 1st paragraph). Thus, the presence of additional nucleoside transporter polypeptides on the surface of bronchial cells might stimulate bronchospasms, by transporting adenosine to the extracellular spaces.

Further research would be required to ascertain the function of SEQ ID NO:1, and to identify a disease with which this polypeptide is associated. Thus, the instant application is an invitation to the skilled artisan to experiment as to the function of the polypeptide of the instant invention and to determine if there is any nexus between said polypeptide any disease or pathological condition.

Utility must be in readily available form. In *Brenner v. Manson*, 148 U.S.P.Q. 689 (Sup. Ct., 1966), a process of producing a novel compound that was structurally analogous to other compounds which were known to possess anti-cancer activity was alleged to be useful because the compound produced thereby was potentially useful as an anti-tumor agent in the absence of evidence supporting this utility. The court expressed the opinion that all chemical compounds are "useful" to the chemical arts when this term is given its broadest interpretation. However, the court held that this

broad interpretation was not the intended definition of "useful" as it appears in 35 U.S.C. § 101, which requires that an invention must have either an immediately obvious or fully disclosed "real world" utility. The instant claims are drawn to a polynucleotide encoding a protein which has undetermined function or biological significance. Until some actual and specific activity can be attributed to the protein identified in the specification as SEQ ID NO:1 or the polynucleotides encoding it (SEQ ID NO:2) the claimed invention is incomplete.

Since the polypeptide of SEQ ID NO:1, or its encoding nucleic acid molecule (SEQ ID NO:2) are not supported by a specific and substantial utility, or a well-established utility, then expression vectors, and transformants comprising the nucleic acids also do not possess utility.

The rejections of Claims 1-12, and 46-48 under 35 U.S.C. 112, first paragraph are maintained for reasons of record. Specifically, since the claimed invention is not supported by either a specific or substantial asserted utility or a well established utility for the reasons of record and those set forth above, one skilled in the art clearly would not know how to use the claimed invention.

Art made of record:

The following art is made of record and not relied upon is considered pertinent to applicant's disclosure. Baker et al. (WO 200012708) teach a polynucleotide (SEQ ID NO:78), encoding a PRO 1380 polypeptide, which has 99.2% identity to SEQ ID NO:2 of the instant invention (See alignment below). However, the reference claims priority to provisional filed 3 November 1998, which is after the perfected priority date of the instant application (27 August 1998)

Human PRO1380 (UNQ717) cDNA sequence SEQ ID NO:78.
WO200012708-A2.
09-MAR-2000.

Art Unit: 1647

Baker K, Goddard A, Gurney AL, Smith V, Watanabe CK, Wood WJ;
Sequence 2243 BP; 463 A; 701 C; 572 G; 507 T; 0 U; 0 Other;

Query Match	99.2%;	Score 2223;	DB 3;	Length 2243;
Best Local Similarity	99.8%;	Pred. No. 0;		
Matches 2226;	Conservative	0;	Mismatches 5;	Indels 0;
			Gaps	0;

Qy	1	CGGCGGCGGTGCGCAGCGGCGACATGGCGTTGTCTCAGAGGACGACTTTCAGCACAGTT	60
Db	13	CGGCGGCGGTGCGCAGCGGCGACATGGCGTTGTCTCAGAGGACGACTTTCAGCACAGTT	72
Qy	61	CAAACTCCACCTACGGAACCAAGCAGCAGTCTCCGAGCTGACGAGGACACTGCTTG	120
Db	73	CAAACTCCACCTACGGAACCAAGCAGCAGTCTCCGAGCTGACGAGGACACTGCTTG	132
Qy	121	AGAAGCTGCTGGACCCGCCGCCCTGGCCTGCAGAGGCCGAGGACGCTTCTGTGGCA	180
Db	133	AGAAGCTGCTGGACCCGCCGCCCTGGCCTGCAGAGGCCGAGGACGCTTCTGTGGCA	192
Qy	181	CATACATCATCTTCTTCAGCCTGGGCATTGGCAGTCTACTGCCATGGAACTCTTTATCA	240
Db	193	CATACATCATCTTCTTCAGCCTGGGCATTGGCAGTCTACTGCCATGGAACTCTTTATCA	252
Qy	241	CTGCCAAGGAGTACTGGATGTTCAAACCTCCGCAACTCCTCCAGCCACGCCACGGGGAGG	300
Db	253	CTGCCAAGGAGTACTGGATGTTCAAACCTCCGCAACTCCTCCAGCCACGCCACGGGGAGG	312
Qy	301	ACCTTGAGGGCTCAGACATCCTGAACTACTTTGAGAGCTACCTTGCGGTGGCTCCACCG	360
Db	313	ACCTTGAGGGCTCAGACATCCTGAACTACTTTGAGAGCTACCTTGCGGTGGCTCCACCG	372
Qy	361	TGCCCTCCATGCTGTGCTGGTGGCCAACTTCCTGCTTGTCAACAGGGTTGCAGTCCACA	420
Db	373	TGCCCTCCATGCTGTGCTGGTGGCCAACTTCCTGCTTGTCAACAGGGTTGCAGTCCACA	432
Qy	421	TCCGTGTCCTGGCCTCACTGACGGTATCCTTGCCCATCTTTCATGGTGATAACTGCATGG	480
Db	433	TCCGTGTCCTGGCCTCACTGACGGTATCCTTGCCCATCTTTCATGGTGATAACTGCATGG	492
Qy	481	TGAAGTGGACACTTCTCCTGGACCGGTGGCTTTTTTGCAGTCAACATTGTCTGCATGG	540
Db	493	TGAAGTGGACACTTCTCCTGGACCGGTGGCTTTTTTGCAGTCAACATTGTCTGCATGG	552
Qy	541	TGATCCTCAGCGGTGCCCTCCACTGTCTTCAGCAGCAGCATCTACGGCATGACCGGCTCCT	600
Db	553	TGATCCTCAGCGGTGCCCTCCACTGTCTTCAGCAGCAGCATCTACGGCATGACCGGCTCCT	612
Qy	601	TTCTATGAGGAACTCCCAAGCACTGATATCAGGAGGACCTGGGCGGACGGTCAAGC	660
Db	613	TTCTATGAGGAACTCCCAAGCACTGATATCAGGAGGACCTGGGCGGACGGTCAAGC	672
Qy	661	CCGTGGCCTCATTGGTGGACTTGGCTGCATCCAGTGATGTGAGGAACAGCGCCTGGCCT	720
Db	673	CCGTGGCCTCATTGGTGGACTTGGCTGCATCCAGTGATGTGAGGAACAGCGCCTGGCCT	732
Qy	721	TCTTCCTGAAGGCAACCATCTTCCTCGTGCCTGCATGGGACTCTACCTGCTGCTGCCA	780
Db	733	TCTTCCTGAAGGCAACCATCTTCCTCGTGCCTGCATGGGACTCTACCTGCTGCTGCCA	792
Qy	781	GGCTGGAGTATGCCAGTACTACATGAGGCCTGTTCTTGCGGCCCATGTGTTTTCTGGTG	840

Art Unit: 1647

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Db      793  GGCTGGAGTATGCCAGGTACTACATGAGGCCTGTTCTTGGCGCCCATGTGTTTCTGGTG 852

Qy      841  AAGAGGAGCTTCCCGAGACTCCCTCAGTGCCCTTCGGTGGCTCCAGATTTCATTGAT 900

Db      853  AAGAGGAGCTTCCCGAGACTCCCTCAGTGCCCTTCGGTGGCTCCAGATTTCATTGAT 912

Qy      901  CCCACACACCCCTCTCCGCCCATCTGAAGAAGACGGCCAGCCTGGGCTTCTGTGTCA 960

Db      913  CCCACACACCCCTCTCCGCCCATCTGAAGAAGACGGCCAGCCTGGGCTTCTGTGTCA 972

Qy      961  CCTACGCTCTTTCATCACAGCCTCATCTACCCCGCGCTGTGCACCAACATCGAGTCC 1020

Db      973  CCTACGCTCTTTCATCACAGCCTCATCTACCCCGCGCTGTGCACCAACATCGAGTCC 1032

Qy      1021  TCAACAAGGGCTCGGGCTCACTGTGGACACCAAGTTTTTCATCCCTCCTCACTACCTTC 1080
|||||
Db      1033  TCAACAAGGGCTCGGGCTCACTGTGGACACCAAGTTTTTCATCCCTCCTCACTACCTTC 1092

Qy      1081  TCCTGTACAACTTTTGCTGACCTATGTGGCCGGCAGCTCACCGCTGGATCCAGGTGCCAG 1140

Db      1093  TCCTGTACAACTTTTGCTGACCTATGTGGCCGGCAGCTCACCGCTGGATCCAGGTGCCAG 1152

Qy      1141  GGCCCAATAGCAAGGCGCTCCCAAGGTTCTGTGCTCCTCCGACCTGCCTCATCCCCCTCT 1200

Db      1153  GGCCCAACAGCAAGGCGCTCCCAAGGTTCTGTGCTCCTCCGACCTGCCTCATCCCCCTCT 1212

Qy      1201  TCGTGCTCTGTAACACAGCCCGCGTCCACCTGAAGACTGTGCTCTTCAGTCCGATG 1260
|||||
Db      1213  TCGTGCTCTGTAACACAGCCCGCGTCCACCTGAAGACTGTGCTCTTCAGTCCGATG 1272

Qy      1261  TGTACCCCGCACTCCTCAGCTCCCTGCTGGGGCTCAGCAACGGCTACCTCAGCACCTTG 1320
|||||
Db      1273  TGTACCCCGCACTCCTCAGCTCCCTGCTGGGGCTCAGCAACGGCTACCTCAGCACCTTG 1332

Qy      1321  CCCTCCTCTACGGGCTTAAGATTGTGCCAGGGAGCTGGCTGAGGCCACGGGAGTGGTGA 1380

Db      1333  CCCTCCTCTACGGGCTTAAGATTGTGCCAGGGAGCTGGCTGAGGCCACGGGAGTGGTGA 1392

Qy      1381  TGTCCCTTTTATGTGTGCTTGGGCTTAACACTGGGCTCAGCCTGCTCTACCCCTCTGGTGC 1440
|||||
Db      1393  TGTCCCTTTTATGTGTGCTTGGGCTTAACACTGGGCTCAGCCTGCTCTACCCCTCTGGTGC 1452

Qy      1441  ACCTCATCTAGAAGGGAGGACACAAGGACATTGGTGCTTCAGAGCCTTTGAAGATGAGAA 1500
|||||
Db      1453  ACCTCATCTAGAAGGGAGGACACAAGGACATTGGTGCTTCAGAGCCTTTGAAGATGAGAA 1512

Qy      1501  GAGAGTCCAGGAGGCTGGGGCCATGGAGGAAGGCCAAAGTTTTCACCTTGGGGACAGA 1560
|||||
Db      1513  GAGAGTCCAGGAGGCTGGGGCCATGGAGGAAGGCCAAAGTTTTCACCTTGGGGACAGA 1572

Qy      1561  GAGCAGAGCACACTCGGGCCTCATCCCTCCCAAGATGCCAGTGAGCCACGTCCATGCCCA 1620
|||||
Db      1573  GAGCAGAGCACACTCGGGCCTCATCCCTCCCAAGATGCCAGTGAGCCACGTCCATGCCCA 1632

Qy      1621  TTCCTGCAAGCGAGATATTCCAGTCATATTACAGAACTCCTGAGACAGTTGAAGAA 1680
|||||
Db      1633  TTCCTGCAAGGGCAGATATTCCAGTCATATTACAGAACTCCTGAGACAGTTGAAGAA 1692

Qy      1681  GAAATAGCACAAATCAGGGGTACTCCCTTCACAGCTGATGGTTAACATTCCCACTTCTTT 1740

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Db      1693  |||||
1693  GAAATAGCACAAATCAGGGGTACTCCCTTCACAGCTGATGGTTAACATTCCACCTTCTTT 1752

Qy      1741  CTAGCCCTTCAAAGATCTGCGCACTGTTTCGCCCTAGAGTTATTACAAAGCCAGTGCCAAA 1800
|||||
Db      1753  CTAGCCCTTCAAAGATGCTGCCAGTGTTCGCCCTAGAGTTATTACAAAGCCAGTGCCAAA 1812

Qy      1801  ACCAGCCATGGGCTCTTTGCAACCTCCCAAGCTGCGCTCATTCAGCTGACAGCGAGATG 1860
|||||
Db      1813  ACCAGCCATGGGCTCTTTGCAACCTCCCAAGCTGCGCTCATTCAGCTGACAGCGAGATG 1872

Qy      1861  CAAGCAATGCTCAGCTCTCCTTACCTGAAGGGGTCTCCCTGGAATGGAAGTCCCTGG 1920
|||||
Db      1873  CAAGCAATGCTCAGCTCTCCTTACCTGAAGGGGTCTCCCTGGAATGGAAGTCCCTGG 1932

Qy      1921  CATGGTCAGTCTTCAGSCCAAGACTCAAGTGTGCACAGACCCCTGTGTCTGTGGGTGA 1980
|||||
Db      1933  CATGGTCAGTCTTCAGSCCAAGACTCAAGTGTGCACAGACCCCTGTGTCTGTGGGTGA 1992

Qy      1981  ACAACTGCCCACTAACAGACTGGAAAAACCCAGAAAGATGGGCCTTCATGAATGCTTCA 2040
|||||
Db      1993  ACAACTGCCCACTAACAGACTGGAAAAACCCAGAAAGATGGGCCTTCATGAATGCTTCA 2052

Qy      2041  TTCCAGAGGGACAGAGGGCCTCCCTGTGCAAGGGATCAAGCATGTCTGGCCTGGGTTT 2100
|||||
Db      2053  TTCCAGAGGGACAGAGGGCCTCCCTGTGCAAGGGATCAAGCATGTCTGGCCTGGGTTT 2112

Qy      2101  CAAAAAAGAGGGATCCTCATGACCTGGTGGTCTATGGCCTGGGTCAAGATGAGGGTCTT 2160
|||||
Db      2113  CAAAAAAGAGGGATCCTCATGACCTGGTGGTCTATGGCCTGGGTCAAGATGAGGGTCTT 2172

Qy      2161  TCAGTGTTCCTGTTTACAACATGTCAAAGCCATTGGTTCAAGGGCGTAATAAATACTTGC 2220
|||||
Db      2173  TCAGTGTTCCTGTTTACAACATGTCAAAGCCATTGGTTCAAGGGCGTAATAAATACTTGC 2232

Qy      2221  GTATTCAAAAA 2231
|||||
Db      2233  GTATTCAAAAA 2243
|||||
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Conclusion:

No claims are allowed.

All claims are drawn to the same invention claimed in the application prior to the entry of the submission under 37 CFR 1.114 and could have been finally rejected on the grounds and art of record in the next Office action if they had been entered in the application prior to entry under 37 CFR 1.114. Accordingly, **THIS ACTION IS MADE FINAL** even though it is a first action after the filing of a request for continued

examination and the submission under 37 CFR 1.114. See MPEP § 706.07(b).
Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Deleted: ¶

Any inquiry concerning this communication or earlier communications from the examiner should be directed to SHULAMITH H. SHAFER whose telephone number is (571)272-3332. The examiner can normally be reached on Monday through Friday, 8 AM to 5 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Manjunath Rao, Ph.D. can be reached on 571-272-0939. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Lorraine Spector/ Ph.D.
Primary Examiner, Art Unit 1647

/S. H. S./
Examiner, Art Unit 1647